



Food and Drug Administration Rockville MD 20857

#10

Re: Differin Solution (5,212,303) (Application 4)

Stephen G. Kunin
Deputy Assistant Commissioner for
Patent Policy and Projects
Office of the Assistant Commissioner for Patents
U.S. Patent and Trademark Office
Crystal Park Building 2, Suite 919
Washington, D.C. 20231

Dear Mr. Kunin:

OCT 24 199

This is in regard to the application for patent term extension for U.S. Patent No. 5,212,303 filed by Centre International de Recherches Dermatologiques ("CIRD") under 35 U.S.C. § 156. The human drug product claimed by the patent is Differin Solution (5,212,303) (Application 4) (adapalene), which was assigned New Drug Application (NDA) No. 20-338.

A review of the Food and Drug Administration's official records indicates that this product was subject to a regulatory review period before its commercial marketing or use, as required under 35 U.S.C. § 156(a)(4). Our records also indicate that it represents the first permitted commercial marketing or use of the product.

The NDA was approved on May 31, 1996, which makes the submission of the patent term extension application on July 26, 1996, timely within the meaning of 35 U.S.C. § 156(d)(1).

Should you conclude that the subject patent is eligible for patent term extension, please advise us accordingly. As required by 35 U.S.C. § 156(d)(2)(A) we will then determine the applicable regulatory review period, publish the determination in the <u>Federal Register</u>, and notify you of our determination.

Please let me know if we can be of further assistance.

Sincerely,

Ronald L. Wilson, Director Health Assessment Policy Staff

Office of Health Affairs

cc: Norman H. Stepno

Burns, Doane, Swecker & Mathis, L.L.P.

P.O. Box 1404

Alexandria, VA 22313-1404





Food and Drug Administration Rockville MD 20857

Re: Differin Topical Gel (5,212,303) (Application 4)

Stephen G. Kunin Deputy Assistant Commissioner for Patent Policy and Projects Office of the Assistant Commissioner for Patents U.S. Patent and Trademark Office Crystal Park Building 2, Suite 919 Washington, D.C. 20231

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A review of the Food and Drug Administration's official records indicates that this product was subject to a regulatory review period before its commercial marketing or use, as required under 35 U.S.C. § 156(a)(4). Our records also indicate that it represents the first permitted commercial marketing or use of the product.

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DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

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